FDA

MAUDE Adverse Event Report: EPIC EPIC CPOE CPOE/EHR COMPUTERIZED PHYSICIAN ENTRY

EPIC EPIC CPOE CPOE/EHR COMPUTERIZED PHYSICIAN ENTRY

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Event Date 10/16/2017
Event Type Injury
Event Description

The report exemplifies widespread (across several hospitals and many pts) toxicity caused by a cpoe (order entry), and ehr (care record) device in which the tests, medications, and treatments that are ordered are not done in a timely manner, if at all. In this case, there are severe obstructive coronary disease. A troponin blood test was ordered to evaluate for myocardial infarction, on more than one occasion. The ancillary service recipient did not carry out the order because it either never got there, was not seen, or other. There is not any reconciliation function to warn that tests and more have not been done. Without knowing that the pt had a myocardial infarction in a timely manner, the correct treatment cannot be carried out. This is potentially life threatening.

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Brand Name EPIC CPOE
Type of Device CPOE/EHR COMPUTERIZED PHYSICIAN ENTRY
Manufacturer *(Section D)* EPIC
Verona WI 53593
MDR Report Key 6964415
Report Number MW5072855
Device Sequence Number 1
Product Code OUG[24]
Report Source Voluntary
Reporter Occupation Physician
Report Date 10/17/2017
***2* DeviceS WERE Involved in the Event:** 1 2
***1* Patient Was Involved in the Event**
Date FDA Received 10/20/2017
Is This An Adverse Event Report? Yes
Is This A Product Problem Report? Yes
Device Operator Health Professional
Was Device Available For Evaluation? Yes
Is The Reporter A Health Professional? Yes
Was the Report Sent to FDA?
Event Location No Information
Was Device Evaluated By Manufacturer?
Is The Device Single Use?
Is this a Reprocessed and Reused Single-Use Device? No
Type of Device Usage

Patient TREATMENT DATA
Date Received: 10/20/2017 Patient Sequence Number: 1

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